

**Table 1.** Eligibility criteria for including studies in this systematic review.

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**Inclusion Criteria:**

1. Study Design: randomised controlled trials and quasi-randomised controlled trials.
2. Types of Participants: reproductive-aged women with a diagnosis of polycystic ovary syndrome (PCOS) based on the National Institute of Health (NIH) diagnostic criteria (1990), the Rotterdam ESHRE/ASRM (2003) diagnostic criteria or the AE-PCOS Criteria (2006). We also included trials where the PCOS diagnosis had been verified by a general practitioner or specialist clinician.
3. Comparators: exercise vs usual care/control, exercise combined with diet vs usual care/control, exercise combined with diet vs diet only. Exercise combined with diet vs exercise only, exercise vs diet, exercise combined with pharmaceutical vs pharmaceutical.
4. All outcomes; Expected outcomes included: primary outcomes, such as blood pressure, fasting blood glucose, insulin and lipid concentrations; and secondary outcomes, such as body mass index, cardiorespiratory fitness, testosterone, free androgen index and health related quality of life measures.

**Exclusion Criteria:**

1. Study Design: case studies, cross sectional and non-randomised controlled trials.
  2. Types of Participants: males, adolescent females, post-menopausal women, women without PCOS
  3. Comparators: women with PCOS vs healthy controls, pharmaceutical vs exercise, pharmaceutical vs diet, diet vs diet, surgical vs any other condition.
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**Table 2.** Characteristics of studies included in this systematic review.

Study (design)	N randomised/ analysed	Intervention Duration (assessment points)	Participant Characteristics (PCOS diagnostic criteria)	Intervention	Outcome measures
Almenning et al. [32] (RCT)	HIIT: 10/8 RT: 11/8 CON: 10/9	10 wks (baseline, 10 wks)	age: 27.2±5.5 y BMI: 26.7±6.0 kg/m <sup>2</sup> (Rotterdam)	HIIT frequency: 3 times/wk HIIT intensity: 2 d/wk, 4 x 4 mins 90-95% HR <sub>max</sub> / 3 x 3 mins ~70% HR <sub>max</sub> . 1 d/wk, 10 x 1 min 'all-out' / 10 x 1 min rest. RT frequency: 3 times/wk RT sets x reps: 3 x 10 RT load: 75% 1-RM	HOMA-IR, FBG, FI, TG, TC, LDL-C, HDL-C VO <sub>2</sub> max, RHR, BW, BMI, WC, BF%, FM, FFM, T, SHBG, FAI, hsCRP.
Brown et al. [96] (RCT)	EX: 21/8 CON: 16/12	20-24 wks due to varying length of ramp up phase (baseline, immediately post)	age: 32.3 ± ns y BMI: 33.0 kg/m <sup>2</sup> (NIH)	Exercise: 12 wk moderate-intensity intervention preceded by 8-12 wk ramp-up. Aerobic duration: ~228 mins/wk (≤ 60 bouts) Aerobic intensity: 40-60% VO <sub>2</sub> max	FBG, FI, HOMA-IR, TG, LDL-C, HDL-C, VO <sub>2</sub> max, BW, BMI, WC, FT, SBP, DBP.
Bruner et al. [97] (RCT)	EX + DIET: 7/7 DIET: 5/5	12 wks (baseline, 12 weeks)	age: 30.7±4.6 y BMI: 36.6±6.0 kg/m <sup>2</sup> (Rotterdam)	Exercise frequency: 3 times/wk Aerobic intensity: 70-85% HR <sub>max</sub> Aerobic duration: 30 mins (+10-min warm-up) RT sets x reps: 2-3 x 10-15 RT load: not specified	FI, QUICKI, VO <sub>2</sub> max, BW, BMI, WC, T, SHBG, FAI
Guzick et al. [98] (RCT)	EX + DIET: 6/6 CON: 6/6	12 wks (baseline, 12 weeks)	age: 31.7±10.0 y BMI: ns (NIH)	Diet: 1 hour/wk of nutritional counselling Exercise frequency: 5 times/wk Exercise intensity: 1050-4200 kJ/wk  Diet: VLCD (8 wks) with calories increased over final 4 wks (4200-5040 kJ/d). 'Optifast' used to supplement diet.	FBG, FI, BW, WHR, T, SHBG, FT, LH, FSH.

Study (design)	N randomised/ analysed	Intervention Duration (assessment points)	Participant Characteristics (PCOS diagnostic criteria)	Intervention	Outcome measures
Hoeger et al. [99] (RCT)	LS + PLA: 11/6 PLA: 9/7 LS + MF : 9/5 MF: 9/5	48 wks (baseline, 24 wks, 48 wks)	age: 28.5±5.2 y BMI: 39.0±6.1 kg/m <sup>2</sup> (NIH)	Exercise programme: Individualised to achieve 150 minutes per wk  Diet: Individualised healthy balanced meal plan to achieve 500-1000 kcal deficit per day  Metformin: 850 mg 2 times/day	BW, T, SHBG, FAI
Konopka et al. [101] (RCT)	EX: 12/12 CON: 13/13	12 wks (baseline, 12 wks)	age: 35±5.0 y BMI: 33.0±5.0 kg/m <sup>2</sup> (Rotterdam)	Exercise frequency: 5 times/wk Exercise intensity: 65% VO <sub>2</sub> peak Exercise duration: 60 mins	FBG, FI, HOMA-IR, BMI, BW, FM, FFM, E <sub>2</sub>
Nasrekani et al. [103] (RCT)	EX: 10/10 CON: 10/10	12 wks (baseline, 12 wks)	age: 30.4±5.9 y BMI: 28.3±6.2 kg/m <sup>2</sup> (Rotterdam)	Exercise frequency: 3 times/wk Exercise intensity: 40-65% HR <sub>max</sub> Exercise duration: 25-30 mins	VO <sub>2</sub> max, BW, BMI, FSH, LH.
Nybacka et al. [104-105] (RCT)	EX: 19/17 EX + DIET: 19/12 DIET: 19/14	4 months (baseline, 4 months)	age: 30.8±5.2 y BMI: 36.0 ± 6.2 kg/m <sup>2</sup> (Rotterdam)	Exercise programme: Individualised to meet individuals' capacity, goals and interest.  Diet: ≥ 600 kcal/day reduction maintaining 55-60% CHO, 25-30% fat and 10-15% protein.	FBG, FI, HOMA-IR, BW, BMI, WHR, BF%, FFM, T, SHBG, FT, E <sub>2</sub> , FSH, LH
Petranyi <i>et al.</i> [106] (QRCT)	LS+MF: 29/29 MF: 27/27	6 months (baseline, 6 months)	Age: 29 ± ns y BMI: 27.2±6.9 kg/m <sup>2</sup> (Rotterdam)	Exercise programme: recommendation to increase physical activity levels. Specifics unclear.  Diet: low glycaemic index diet with caloric restriction for those who are obese.  Metformin: 500 mg 3 times/day	BMI, WHR

Study (design)	N randomised/ analysed	Intervention Duration (assessment points)	Participant Characteristics (PCOS diagnostic criteria)	Intervention	Outcome measures
Roessler et al. [34] (Randomised crossover)	EX: 8/7 CON: 9/7	16 wks (baseline, 8 wks, 16 wks)	age: 31.7±7.9 y BMI: 36.3±7.2 kg/m <sup>2</sup> (Rotterdam)	Exercise frequency: 3 times/wk (2 x cycle, 1 x walk) Exercise intensity: following 2-week ramp, cycling 20-180 secs 80-100% HR <sub>max</sub> / rest 25-180 secs 45-65% HR <sub>max</sub> . Walking 3-5 mins 80-90% HR <sub>max</sub> / 1 min 50-60% HR <sub>max</sub> . Exercise duration: 45 mins (+10 min warm-up). Control: Group counselling sessions (2 hours, 1 time/wk) focussing on barriers and motivation.	VO <sub>2</sub> max, BW, BMI, WC
Sa et al. [107-108] (RCT)	EX: 15/14 CON: 15/13	16 wks (baseline, 16 wks)	age: 26.0±5.0 y BMI: 32.8±4.6 kg/m <sup>2</sup> (Rotterdam)	Exercise frequency: 3 times/wk Exercise intensity: 60-85% HR <sub>max</sub> Exercise duration: 40 mins (+5 mins)	SBP, DBP, FI, BMI, RHR, VO <sub>2</sub> max, T, FSH, LH
Saremi et al. [109] (RCT)	EX: 11/11 CON: 11/11	8 wks (baseline, 8wks)	age: 35.2±4.4 y BMI: 28.3±4.3 kg/m <sup>2</sup> (Rotterdam)	Exercise frequency: 3 times/wk Exercise intensity: 40-65% HR <sub>max</sub> Exercise duration: 30 mins	FBG, FI, HOMA-IR, TG, TC, LDL-C, HDL-C, VO <sub>2</sub> peak, BW, BMI, BF%, WC, WHR
Saremi et al. [110] (RCT)	EX + PLA: 10/10 CON : 10/10 EX + CAL: 10/10	8 wks (baseline, 8 wks)	age: 27.1±5.1 y BMI: 25.5±2.7 kg/m <sup>2</sup> (Rotterdam)	Exercise frequency: 3 times/wk RT sets x reps: 1-2 x 15-20 RT load: 40-60% 1-RM	FBG, FI, HOMA-IR, TG, TC, LDL-C, HDL-C, BW, BMI
Stener-Victorin et al. [100, 102, 111-113] (RCT)	EX: 34/22 CON: 17/13 ACU: 33/24	16 wks (baseline, 16 wks, 32 wks)	age: 30±4.4 y BMI: 28.1±7.3 kg/m <sup>2</sup> (Rotterdam)	Exercise frequency: 3 times/wk Exercise intensity: HR ≥ 120 BPM Exercise duration: 30-45 mins  Low-frequency electroacupuncture: 14 x 30 min treatments over 16 wks.	SBP, DBP, FBG, FI, HOMA-IR, TG, TC, LDL-C, HDL-C, BMI, WHR, T, FT, SHBG, FAI, LH, FSH, VO <sub>2</sub> max, BMI, E <sub>2</sub>

Study (design)	N randomised/ analysed	Intervention Duration (assessment points)	Participant Characteristics (PCOS diagnostic criteria)	Intervention	Outcome measures
Thomson et al. [33, 114-116] (RCT)	AET + DIET: 31/18 AET + RT + DIET: 33/20 DIET: 30/14	20 weeks (baseline, 10 wks, 20 wks)	age: 29.3±6.8 y BMI: 36.1±4.8 kg/m <sup>2</sup> (Rotterdam)	Exercise frequency: 5 times/wk (3 x aerobic, 2 x RT in combined exercise group) Aerobic intensity: 60-65% HR <sub>max</sub> progressed to 75-80% HR <sub>max</sub> by study end Aerobic duration: 25-30 mins progressed to 45 mins by study end RT sets x reps: 3 x 12 RT load: 50-60% 1-RM progressed to 65-75% 1-RM after 2 weeks Diet: energy restricted high protein diet (5000-6000 kJ/day).	SBP, DBP, FBG, FI, HOMA-IR, TG, TC, LDL-C, HDL-C, BW, BF%, FM, FFM, WC, T, SHBG, FAI, PCOS-Q
Turan et al. [117] (RCT)	EX: 16/14 CON: 16/16	8 wks (baseline, 8 wks)	age: 24.5 ± 2.8 y BMI: 21.9±3.5 kg/m <sup>2</sup> (Rotterdam)	Exercise frequency: 3 times/wk Exercise duration: 50-60 mins Aerobic intensity: 65-70% HR <sub>max</sub> RT sets x reps: 1 x 15 RT load: 5-6 on RPE for RT scale	SBP, DBP, FBG, HOMA-IR, FI, TG, TC, HDL-C, LDL-C, BMI, WC, RHR, VO <sub>2</sub> max, T, FT, E <sub>2</sub> , LH, FSH
Vigorito et al. [118] (RCT)	EX: 45/45 CON: 45/45	3 months (baseline, 3 months)	age: 21.8±2.1 y BMI: 29.4±3.2 kg/m <sup>2</sup> (Rotterdam)	Exercise frequency: 3 times/wk Exercise intensity: 60-70% VO <sub>2</sub> max Exercise duration: 30 mins	SBP, DBP, FBG, FI, TG, TC, LDL-C, HDL-C, VO <sub>2</sub> max, RHR, BMI, WC, E <sub>2</sub> , T, FT, SHBG, FAI, LH, FSH, CRP
Vizza et al. [119] (RCT)	EX: 8/7 CON: 7/6	12 wks (baseline, 12 wks)	age: 27±5.0 y BMI: 37.8±11.4 kg/m <sup>2</sup>	Exercise frequency: 4 times/wk (2 x RT, 2 home-based) RT sets x reps: 2-3 x 8-12 RT load: Progressed with strength gains Home-based: Callisthenics, 3 sets of 10 reps	FBG, FI, HOMA-IR, BW, BMI, WC, FM, FFM, BF%, hsCRP, T, SHBG, FAI, PCOS-Q, SF-36

Studies presented by lead author and year of publication. Design; RCT: randomised controlled trial, QRCT: quasi-randomised controlled trial. N randomised: the number of participants randomised into each study arm at the study initiation; analysed is the number of participants included within the analysis; HIIT: high-intensity interval training; RT: resistance training; CON: control group; EX: exercise group; DIET: dietary intervention; LS: lifestyle; PLA: placebo; MF: Metformin; ACU: acupuncture; AET: aerobic

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exercise training; CAL: calcium supplementation. Intervention duration: length of the duration; assessment points: the time-points at which researchers have assessed outcome measures. Participant characteristics presented as mean  $\pm$  standard deviation (SD) or median in one study [96] for age (in years; y) and BMI ( $\text{kg}/\text{m}^2$ ) at study entry; ns: not specified. Diagnostic criteria: the specific criteria used to confirm a PCOS diagnosis; NIH: National Institute of Health (1990) diagnostic criteria; Rotterdam: European Society for Human Reproductive and Embryology/American Society for Reproductive Medicine (2003). Outcome measures refers to the outcomes from each study that are relevant to this systematic review.  $\text{VO}_2$  max: maximum oxygen uptake; RHR: resting heart rate; HDL-C: high density lipoprotein cholesterol; LDL-C: low density lipoprotein cholesterol; TC: total cholesterol; TG: triglycerides; FBG: fasting blood glucose; FI: fasting insulin; HOMA-IR: homeostatic assessment of insulin resistance; QUICKI: quantitative insulin sensitivity check index; FM: fat mass; FFM: fat free mass; BF%: body fat percentage; BW: body weight; BMI: body mass index; WC: waist circumference; WHR: waist to hip ratio; SHBG: sex hormone binding globulin; FAI: free androgen index; T: testosterone; FT: free testosterone;  $\text{E}_2$ : oestradiol; LH: luteinising hormone; FSH: follicle stimulating hormone; SBP: systolic blood pressure; DBP: diastolic blood pressure; hsCRP: high-sensitivity C-reactive protein; d: day; mins: minutes; wk: week; reps: repetitions; RM: maximum number of repetitions; HRmax: maximum heart rate; PCOSQ: PCOS health-related questionnaire; SF-36: Optum36-item Short Form Survey; VLCD: very low calorie diet; CHO: carbohydrate.

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**Table 3.** Effect estimates and heterogeneity for change from baseline to post-intervention scores and immediately post-intervention values, for all outcomes analysed in the exercise versus control comparison

Outcome	Change from baseline						Immediately post-intervention values					
	Trials	N	MD	Lower 95% CI	Upper 95% CI	I <sup>2</sup> (%)	Trials	N	MD	Lower 95% CI	Upper 95% CI	I <sup>2</sup> (%)
SBP (mmHg)	4	158	-2.93	-7.06	1.20	50	4	158	2.02	-6.82	10.86	87
DBP (mmHg)	4	158	-2.19	-5.23	0.85	46	4	158	-0.82	-3.49	1.84	31
FBG (mg/dL)	9	263	-1.08	-2.47	0.30	16	8	238	-1.69	-4.35	0.97	37
FI (μIU/mL)	9	263	-2.44**	-4.24	-0.64	91	8	238	-2.11**	-3.49	-0.73	40
HOMA-IR	8	173	-0.57**	-0.99	-0.14	87	7	148	-0.22	-0.80	0.36	69
Total Cholesterol (mg/dL)	7	225	-5.88**	-9.92	-1.83	35	7	225	-6.35**	-10.76	-1.95	0
LDL-C (mg/dL)	7	225	-7.39***	-9.83	-4.95	0	7	225	-6.68**	-11.66	-1.70	0
HDL-C (mg/dL) ▲	7	225	0.29	-1.46	2.04	52	7	225	1.87	-1.59	5.33	65
Triglycerides (mg/dL)	7	225	-4.78***	-7.52	-2.05	3	7	225	-1.97	-7.36	3.42	18
VO <sub>2</sub> max (ml/kg/min)	6	229	3.84***	2.87	4.81	17	5	184	5.01***	3.48	6.54	42
Resting Heart Rate (bpm)	4	156	-2.65	-5.55	0.25	51	4	156	-3.26***	-4.93	-1.59	0
BMI (kg/m <sup>2</sup> )	11	331	-0.49	-1.04	0.06	66	10	272	-1.02**	-1.81	-0.23	0
Body Mass (kg)	7	139	-1.25	-3.27	0.76	33	6	128	-0.48	-4.86	3.91	0
WC (cm)	7	221	-2.62***	-4.13	-1.11	53	7	221	-2.33	-5.23	0.58	15
WHR	2	101	-0.03	-0.08	0.02	0	2	101	-0.04	-0.08	0.01	19
Body Fat (%)	3	60	-1.39*	-2.61	-0.18	30	3	60	-3.28	-7.39	0.83	22
Fat Mass (kg)	3	63	-1.70	-3.93	0.53	70	2	38	5.14	-14.39	24.68	65
FFM (kg)	3	63	0.46	-0.89	1.81	58	2	38	4.99	-7.31	17.28	75
Testosterone (nmol/L)	5	203	-0.09	-0.24	0.06	0	5	169	-0.08	-0.35	0.19	37
SHBG (nmol/L)	4	173	7.51	-8.01	23.04	89	4	139	4.03	-18.57	26.63	66
Free Testosterone (pg/mL)	2	74	-0.43	-1.74	0.88	76	2	41	0.33	-0.10	0.77	0
FAI	4	139	0.24	-0.55	1.04	0	4	139	0.68	-1.09	2.44	46
FG	2	135	-0.63	-2.08	0.81	0	2	101	-0.75	-2.03	0.54	0
Oestradiol (pmol/L)	4	190	-13.94	-54.53	26.64	65	2	120	0.27	-11.27	11.80	0
DHEA-S (μmol/L)	2	70	-0.60	-1.58	0.39	0	2	36	-0.20	-1.87	1.46	0
LH (IU/L)	4	185	-0.30	-2.54	1.95	72	4	151	-0.66	-2.39	1.06	43
FSH (IU/L)	4	185	0.23	-0.08	0.53	0	4	151	-0.01	-0.40	0.37	0
LH/FSH ratio	2	41	-0.02	-0.38	0.33	0	2	41	0.32	-0.22	0.86	37
Progesterone (nmol/L)	2	115	-0.72	-2.53	1.09	74	-	-	-	-	-	-
Prolactin (ng/mL)	2	110	-0.05	-0.71	0.61	0	2	110	0.20	-0.27	0.68	0
hsCRP (mg/L)	2	38	-0.41	-1.19	0.37	0	2	38	0.67	-1.31	2.65	0
AMH (ng/mL)	3	67	-0.67	-1.65	0.32	0	2	67	0.48	-1.89	2.84	0

Adiponectin ( $\mu\text{g/mL}$ )	2	70	-0.20	-1.04	0.64	0	-	-	-	-	-	-
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**Key:** 95% CI: 95% confidence intervals; SBP: systolic blood pressure; DBP: diastolic blood pressure; FBG: fasting blood glucose; FI: fasting insulin; HOMA-IR: homeostatic model of assessment - insulin resistance; LDL-C: low-density lipoprotein cholesterol; HDL-C: high-density lipoprotein cholesterol; BMI: body mass index; WC: waist circumference; WHR: waist to hip ratio; FFM: fat free mass; SHBG: sex hormone binding globulin; FAI: free androgen index; FG: Ferriman-Gallwey score; DHEA-S: dehydroepiandrosterone sulfate; LH: luteinising hormone; FSH: follicle stimulating hormone; hsCRP: high-sensitivity C-reactive protein; AMH: anti-Müllerian hormone. ▲: positive values favour exercise over control. Statistically significant effects denoted by: \*  $P \leq 0.05$ ; \*\*  $P \leq 0.01$ ; \*\*\*  $P \leq 0.001$ . Trials: number of studies included within analysis,  $N$ : number of participants included within analysis. Effect estimates are reported as mean differences (MD), and 95% confidence intervals, between exercise and usual care groups. Heterogeneity reported using  $I^2$  statistic.

**Table 4.** Summary of findings for Primary Outcomes: exercise versus control.

<b>Exercise compared to usual care for women with PCOS</b>						
<b>Patient or population:</b> women with PCOS						
<b>Setting:</b>						
<b>Intervention:</b> exercise						
<b>Comparison:</b> usual care						
Outcomes	Anticipated absolute effects* (95% CI)		Relative effect (95% CI)	№ of participants (studies)	Certainty of the evidence (GRADE)	Comments
	Risk with usual care	Risk with exercise				
Systolic blood pressure (change from baseline) follow up: range 8 weeks to 16 weeks	The mean systolic blood pressure (change from baseline) ranged from <b>-2.5 to 1.1</b> mmHg	The mean systolic blood pressure (change from baseline) in the intervention group was 2.93 mmHg lower (7.06 lower to 1.2 higher)	-	158 (4 RCTs)	⊕⊕○○ LOW <sup>a,b</sup>	Exercise may result in little to no difference in systolic blood pressure (change from baseline).
Diastolic blood pressure (change from baseline) follow up: range 8 weeks to 16 weeks	The mean diastolic blood pressure (change from baseline) ranged from <b>-3.1 to 2.9</b> mmHg	The mean diastolic blood pressure (change from baseline) in the intervention group was 2.19 mmHg lower (5.23 lower to 0.85 higher)	-	158 (4 RCTs)	⊕⊕○○ LOW <sup>a,b</sup>	Exercise may result in little to no difference in diastolic blood pressure (change from baseline).
Fasting blood glucose (change from baseline) follow up: range 8 weeks to 16 weeks	The mean fasting blood glucose (change from baseline) ranged from <b>-1.3 to 2.6</b> mg/dL	The mean fasting blood glucose (change from baseline) in the intervention group was 1.08 mg/dL lower (2.47 lower to 0.3 higher)	-	263 (9 RCTs)	⊕⊕○○ LOW <sup>c,d</sup>	Exercise may result in little to no difference in fasting blood glucose (change from baseline).

**Table 4.** Summary of findings for Primary Outcomes: exercise versus control.

<b>Exercise compared to usual care for women with PCOS</b>						
<b>Patient or population:</b> women with PCOS						
<b>Setting:</b>						
<b>Intervention:</b> exercise						
<b>Comparison:</b> usual care						
Outcomes	Anticipated absolute effects* (95% CI)		Relative effect (95% CI)	№ of participants (studies)	Certainty of the evidence (GRADE)	Comments
	Risk with usual care	Risk with exercise				
Fasting insulin (change from baseline) follow up: range 8 weeks to 16 weeks	The mean fasting insulin (change from baseline) ranged from <b>-4.1 to 2.5</b> µU/ml	The mean fasting insulin (change from baseline) in the intervention group was 2.44 µU/ml lower (4.42 lower to 0.64 lower)	-	263 (9 RCTs)	⊕○○○ VERY LOW <sup>e,f,g</sup>	Exercise may reduce fasting insulin (change from baseline) but we are very uncertain.
HOMA-IR (change from baseline) follow up: range 8 weeks to 16 weeks	The mean HOMA-IR (change from baseline) ranged from <b>-0.4 to 0.7</b>	The mean HOMA-IR (change from baseline) in the intervention group was 0.57 lower (0.99 lower to 0.14 lower)	-	173 (8 RCTs)	⊕○○○ VERY LOW <sup>d,e,h</sup>	Exercise may reduce HOMA-IR (change from baseline) but we are very uncertain.
Total cholesterol (change from baseline) follow up: range 8 weeks to 16 weeks	The mean total cholesterol (change from baseline) ranged from <b>-8.85 to 6.85</b> mg/dL	The mean total cholesterol (change from baseline) in the intervention group was 6.48 mg/dL lower (10.5 lower to 2.45 lower)	-	225 (7 RCTs)	⊕⊕○○ LOW <sup>g,i</sup>	Exercise may reduce total cholesterol (change from baseline) slightly.

**Table 4.** Summary of findings for Primary Outcomes: exercise versus control.

<b>Exercise compared to usual care for women with PCOS</b>						
<b>Patient or population:</b> women with PCOS						
<b>Setting:</b>						
<b>Intervention:</b> exercise						
<b>Comparison:</b> usual care						
Outcomes	Anticipated absolute effects* (95% CI)		Relative effect (95% CI)	№ of participants (studies)	Certainty of the evidence (GRADE)	Comments
	Risk with usual care	Risk with exercise				
LDL-C (change from baseline) follow up: range 8 weeks to 16 weeks	The mean LDL-C (change from baseline) ranged from <b>-17.7 to 7.03</b> mg/dL	The mean LDL-C (change from baseline) in the intervention group was 7.51 mg/dL lower (10.01 lower to 5.02 lower)	-	225 (7 RCTs)	⊕⊕○○ LOW <sup>g,i</sup>	Exercise may reduce LDL-C (change from baseline) slightly.
HDL-C (change from baseline) follow up: range 8 weeks to 16 weeks	The mean HDL-C (change from baseline) ranged from <b>-17.7 to 3.5</b> mg/dL	The mean HDL-C (change from baseline) in the intervention group was 0.01 mg/dL lower (1.91 lower to 1.89 higher)	-	225 (7 RCTs)	⊕⊕○○ LOW <sup>g,i</sup>	Exercise may result in little to no difference in HDL-C (change from baseline).
Triglycerides (change from baseline) follow up: range 8 weeks to 16 weeks	The mean triglycerides (change from baseline) ranged from <b>-1.0 to 8.9</b> mg/dL	The mean triglycerides (change from baseline) in the intervention group was 4.78 mg/dL lower (7.52 lower to 2.05 lower)	-	225 (7 RCTs)	⊕⊕○○ LOW <sup>g,i</sup>	Exercise likely results in a small effect that may not be an important (or unimportant) reduction in triglycerides (change from baseline).

\*The risk in the intervention group (and its 95% confidence interval) is based on the assumed risk in the comparison group and the **relative effect** of the intervention (and its 95% CI).

CI: Confidence interval; MD: Mean difference

**Table 4.** Summary of findings for Primary Outcomes: exercise versus control.

**Exercise compared to usual care for women with PCOS**

**Patient or population:** women with PCOS

**Setting:**

**Intervention:** exercise

**Comparison:** usual care

Outcomes	Anticipated absolute effects* (95% CI)		Relative effect (95% CI)	№ of participants (studies)	Certainty of the evidence (GRADE)	Comments
	Risk with usual care	Risk with exercise				

**GRADE Working Group grades of evidence**

**High certainty:** We are very confident that the true effect lies close to that of the estimate of the effect

**Moderate certainty:** We are moderately confident in the effect estimate: The true effect is likely to be close to the estimate of the effect, but there is a possibility that it is substantially different

**Low certainty:** Our confidence in the effect estimate is limited: The true effect may be substantially different from the estimate of the effect

**Very low certainty:** We have very little confidence in the effect estimate: The true effect is likely to be substantially different from the estimate of effect

**Explanations**

- a. Three of the four trials had a high or unclear risk of selection bias, detection bias, and reporting bias; all were at high risk of performance bias; two were at high or unclear risk of attrition bias; and all were at a high or unclear risk of contamination. Therefore we downgraded by one level.
- b. Small number of participants, wide confidence intervals for three of the four trials, and null/negligible effect and appreciable benefit included in the confidence interval for the mean difference. Therefore, we downgraded by one level.
- c. Most trials were at an unclear or high risk of selection bias, detection bias, and reporting bias; and all trials were at a high or unclear risk of contamination and low adherence. Therefore, we downgraded by one level.
- d. Small number of participants and null/negligible effect and appreciable benefit included in the confidence interval for the mean difference. Therefore, we downgraded by one level.
- e. Most trials were at an unclear or high risk of selection bias, detection bias, attrition bias, and reporting bias; and most trials were at a high or unclear risk of contamination and low adherence. Therefore, we downgraded by one level.
- f. Considerable heterogeneity was observed. Therefore, we downgraded by one level.
- g. Small number of participants and wide confidence intervals in the included trials. Therefore, we downgraded by one level.
- h. Considerable heterogeneity was observed and there was minimal or no overlap of confidence intervals. Therefore, we downgraded by one level.
- i. Most trials were at an unclear or high risk of selection bias, detection bias, and reporting bias; and all trials were at a high or unclear risk of contamination. Therefore, we downgraded by one level.

**Table 5.** Summary of findings for Primary Outcomes: exercise and diet versus diet.

<b>Exercise and diet compared to Diet for women with PCOS</b>						
Patient or population: women with PCOS						
Setting:						
Intervention: exercise and diet						
Comparison: Diet						
Outcomes	Anticipated absolute effects* (95% CI)		Relative effect (95% CI)	No of participants (studies)	Certainty of the evidence (GRADE)	Comments
	Risk with Diet	Risk with exercise and diet				
Fasting blood glucose (change from baseline) follow up: range 16 weeks to 20 weeks	The mean fasting blood glucose (change from baseline) ranged from <b>-7.0 to -3.2</b> mg/dL	The mean fasting blood glucose (change from baseline) in the intervention group was 2.92 mg/dL higher (0.4 lower to 6.23 higher)	-	78 (2 RCTs)	⊕○○○ VERY LOW <sup>a,b</sup>	We are uncertain about the effect of exercise and diet on fasting blood glucose (change from baseline).
Fasting insulin (change from baseline) follow up: range 12 weeks to 20 weeks	The mean fasting insulin (change from baseline) ranged from <b>-2.9 to -18.54</b> μU/ml	The mean fasting insulin (change from baseline) in the intervention group was 2.22 μU/ml higher (3.7 lower to 8.14 higher)	-	90 (3 RCTs)	⊕○○○ VERY LOW <sup>a,c,d</sup>	We are uncertain about the effect of exercise and diet on fasting insulin (change from baseline).
HOMA-IR (change from baseline) follow up: range 16 weeks to 20 weeks	The mean HOMA-IR (change from baseline) ranged from <b>-0.74 to -0.56</b>	The mean HOMA-IR (change from baseline) in the intervention group was 0.01 lower (0.45 lower to 0.43 higher)	-	78 (2 RCTs)	⊕○○○ VERY LOW <sup>a,b</sup>	We are uncertain about the effect of exercise and diet on HOMA-IR (change from baseline).

\*The risk in the intervention group (and its 95% confidence interval) is based on the assumed risk in the comparison group and the **relative effect** of the intervention (and its 95% CI).

CI: Confidence interval; MD: Mean difference

**Table 5.** Summary of findings for Primary Outcomes: exercise and diet versus diet.

Exercise and diet compared to Diet for women with PCOS						
Patient or population: women with PCOS						
Setting:						
Intervention: exercise and diet						
Comparison: Diet						
Outcomes	Anticipated absolute effects* (95% CI)		Relative effect (95% CI)	No of participants (studies)	Certainty of the evidence (GRADE)	Comments
	Risk with Diet	Risk with exercise and diet				

**GRADE Working Group grades of evidence**

**High certainty:** We are very confident that the true effect lies close to that of the estimate of the effect

**Moderate certainty:** We are moderately confident in the effect estimate: The true effect is likely to be close to the estimate of the effect, but there is a possibility that it is substantially different

**Low certainty:** Our confidence in the effect estimate is limited: The true effect may be substantially different from the estimate of the effect

**Very low certainty:** We have very little confidence in the effect estimate: The true effect is likely to be substantially different from the estimate of effect

**Explanations**

- a. All trials were at an unclear risk of selection bias, reporting bias, contamination, and adherence issues. All trials were at a high risk of detection bias and attrition bias. Therefore, we downgraded by one level.
- b. Small number of participants, only two trials, and wide confidence intervals in the included trials. Therefore, we downgraded by two levels.
- c. Substantial heterogeneity was observed. Therefore, we downgraded by one level.
- d. Small number of participants and trials, wide confidence intervals, and null/negligible effect and appreciable benefit included in the confidence interval for the mean difference. Therefore, we downgraded by two levels.

